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(54) Title: CORROSION INHIBITING PROCESSES FOR REFRIGERATION SYSTEMS

(57) Abstract

Organic compounds useful as anticorrosion agents in refrigeration systems. The organic compounds can be added directly into the refrigeration systems. Alternatively the organic compound can be added to an alkali metal halide absorption refrigeration solutions. The organic compounds can minimize corrosion of the refrigeration system.

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CORROSION INHIBITING PROCESSES FOR REFRIGERATION SYSTEMS

Field of the Invention

This invention relates generally to minimizing corrosion in refrigeration systems, and in particular to processes and solutions which include 5 organic corrosion inhibiting agents for absorption refrigeration systems.

Background of the Invention

Absorption refrigeration machines are widely used in commercial operations. A typical absorption 10 refrigeration machine includes four major sections: absorber, generator, condenser, and evaporator.

In the absorber section, an absorbent fluid, typically an aqueous alkali metal halide solution, such as a lithium bromide solution, absorbs a refrigerant, 15 typically water vapor. The absorber can be operated under sub-atmospheric pressure. The resultant weak or diluted absorbent fluid (about 40-58% concentration of alkali metal halide rich in the refrigerant) is pumped to the generator. Here heat is applied to the fluid to 20 partially boil off the refrigerant to again concentrate the absorbent fluid. The concentrated absorbent solution (now about 63-65% alkali metal halide) from the generator is passed through a heat exchanger and then sprayed back into the absorber section where it 25 resumes absorption of the refrigerant vapor.

The refrigerant vapor liberated in the generator migrates to the condenser where it is liquefied by exchanging heat with a cooling fluid (typically water) flowing through tubing (typically 30 copper based alloy tubes) present in the condenser section. The liquid refrigerant in the condenser moves into the evaporator, which is also operated at an extremely low pressure. In the evaporator, the

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refrigerant cools the relatively warm system water circulating through the tubes of the tube bundle, and the chilled water is circulated to the load. Heat from the system water vaporizes the refrigerant water which 5 then migrates to the absorber section for absorption into the concentrated solution and completes the cycle.

Aqueous alkali metal halide solutions are widely used as absorption fluids in commercial absorption refrigeration systems. An exemplary alkali 10 metal halide solution for this application is a lithium bromide solution, adjusted to pH range of 7-13 with lithium hydroxide. Although this and other types of absorption solutions can be advantageous for the refrigeration cycles, alkali metal halides can be 15 corrosive towards the materials used to construct the refrigeration machine. Such materials can include mild and stainless steel for containment components and copper or copper-nickel alloys for tube bundles, among others.

20 In addition to the surface damage caused by corrosion, the corrosion reaction evolves hydrogen gas as a byproduct. Incondensibles in the form of atoms or ions can easily enter and diffuse into metals, resulting in the degradation of their mechanical 25 properties under certain system conditions. Incondensible gases also affect the performance of the machine.

The severity of corrosion can vary, depending upon factors such as temperature of the system, 30 concentration of alkali metal halide in the absorption solution, metals used in the construction of the refrigeration unit, the presence of air, and the like. For example, during use, the internal temperatures of such machines can be high, typically up to about 450°F 35 and higher, depending on the type of the absorption cycle, which can increase the corrosive effect of the alkali metal halide solution.

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Various additives, such as lithium chromate, lithium nitrate, and lithium molybdate, have been proposed as corrosion inhibitors in alkali metal halide absorption solutions. However, lithium chromate can raise environmental concerns, and its use is being phased out. Further, the level of chromate and its oxidation state must be carefully maintained. If too little chromate is used, then it does not properly passivate the whole metal surface and pitting can result. Lithium nitrate can potentially evolve ammonia, which can cause stress corrosion cracking of copper based alloys such as heat exchanger tubes. Lithium molybdate exhibits only limited solubility in alkali metal halide solutions. In addition, lithium molybdate is metastable in aqueous halide solutions and thus it can be difficult to maintain a constant concentration of molybdate ions in solution.

Summary of the Invention

The present invention provides processes for inhibiting corrosion of refrigeration systems which use an absorbent fluid, typically an aqueous alkali metal halide solution, such as a lithium bromide solution. In the invention, a suitable organic compound can be added directly into the refrigeration machine. Alternatively, the organic compound can be added to an absorbent fluid, and the fluid then added to and circulated through the refrigeration system.

The use of organic compounds as anticorrosion agents can provide comparable and even improved corrosion inhibition as compared to solutions which include conventional corrosion inhibition additives. For example, the invention can provide good anticorrosion benefits without pitting and severe crevice corrosion which can result for carbon steel at high temperatures using conventional corrosion inhibitors such as lithium chromate.

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Other advantages include reduced corrosion of the construction materials of the refrigeration machine, such as carbon steel, due to vapor phase, reduced amount of hydrogen generation up to 400°F and
5 enhanced performance of the chillers, due to reduced accumulation of non-condensable gas in the absorber.

Still further, use of the organic compounds in a refrigeration machine can result in a protective layer of magnetite (iron oxide) formed on carbon steel.
10 No pitting or crevice corrosion were found in the presence of additives disclosed in this invention.

Brief Description of the Drawings

Some of the features and advantages of the invention having been described, others will become
15 apparent from the detailed description which follows, and from the accompanying drawings, in which:

Figure 1 is an x-ray diffraction pattern of a surface layer formed on carbon steel coupons exposed to 8-hydroxyquinoline (1000 ppm)/LiBr solution (65 wt.%)
20 at 400°F for one week; and

Figure 2 is a graph illustrating surface analysis of the coupon of Figure 1 using scanning electron microscopy/energy dispersive spectrometry (SEM/EDS) technique.

Detailed Description of the Invention

Organic compounds useful as corrosion inhibiting agents in refrigeration systems which use an alkali metal halide absorption solution are selected from those organics which can exhibit corrosion
30 inhibiting properties in absorption refrigeration systems. Exemplary organic compounds useful in the invention include, but are not limited to, heterobicyclic organic compounds which contain at least one nitrogen atom as part of the ring structure; thio-carboxylic acids containing heterocyclic substituents;
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organophosphates; and mixtures thereof. Exemplary nitrogen-containing heterobicyclic organic compounds include, but are not limited to, quinoline, tetrahydroquinoline, isoquinoline, 5 tetrahydroisoquinoline, quinazoline, quinoxaline, 2,3-benzopyrrole, carboxylic, aldehyde or hydroxy derivatives thereof, and mixtures thereof.

Exemplary thio-carboxylic acids containing heterocyclic substituents include benzothiazoyl-thio-carboxylic acids, benzooxazoyl-thio-carboxylic acids, and benzimidazoyl-thio-carboxylic acids, such as (2-benzothiazoylylthio)-butanedioic acid, commercially available from Ciba-Giegy Corporation as "Irgacor", (2-benzothiazoylylthio)-succinic acid, and the like and mixtures thereof. These and others are described, for example, in U.S. Patent No. 2,725,364; DE 19623268; U.S. Patent No. 4,612,378; U.S. Patent No. 5,152,929; and U.S. Patent No. 5,347,008, the entire disclosure of each of which is hereby incorporated by reference.

Exemplary organophosphates include, but are not limited to, substituted and unsubstituted triaryl phosphates, such as trixylenyl phosphate, commercially available from FMC Corporation as "Durad 220X" and tricresyl phosphate, commercially available from FMC Corporation as "Durad 125"; alkylated phenyl phosphate esters and alkyl acid orthophosphates, such as that commercially available from Octel America Inc. as "Ortholeum", as well as mixtures thereof. These compounds are described, for example, in U.S. Patent Nos. 3,424,601 and 5,206,404, the entire disclosure of each of which is hereby incorporated by reference.

Although not wishing to be bound by any explanation of the invention, it is currently believed that the organic compounds can be absorbed onto surfaces present in the refrigeration system and can form a protective layer thereon. Typically, organic compounds have limited solubility in alkali metal

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halide solutions, such as lithium bromide solutions. This has limited the use of such compounds in absorption refrigeration solutions. However, the inventors have found that the organic compounds are 5 capable of providing protection so long as the compound is introduced into the refrigeration system in quantities sufficient to provide a protective, anticorrosion effect. For example, a suitable organic compound can be added directly into the refrigeration 10 machine. Alternatively, the organic compound can be added to the absorbent fluid, and the fluid then circulated through the refrigeration system. Even organic compounds with little or no solubility in brine solutions can be circulated through the system and 15 provide a protective effect. Further, various organic compounds do exhibit some solubility in the brine absorption solutions at increased temperatures. In this regard, the brine solution can be preheated and organic compound added to the heated solution prior to 20 addition to the refrigeration system. Even if the organic compound is added directly into the refrigeration system, at least some of the organic compound can be solubilized in the brine solution due to the high operating temperatures of the system.

25 The organic compounds can be used singly or as mixtures with one another and/or with other corrosion inhibiting agents. The organic compounds are present in the refrigeration system in amounts sufficient to provide the desired corrosion inhibiting 30 effect. This amount can vary depending upon various factors, such as the solubility of the organic compound in the absorption solution, the nature of organic compound, temperature of the environment of the refrigeration machine, concentration of the alkali 35 metal halide solution, metals used in the construction of the refrigeration unit, the presence of air, and the like. Preferably, organic compounds are added to

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refrigeration systems in an amount ranging from about 100 ppm to about 5000 ppm, and more preferably from about 300 to about 3000 ppm.

The absorption solutions include alkali metal halide in conventional amounts. An exemplary alkali metal halide solution includes alkali metal halide in an amount from about 40 to about 65 weight percent, preferably about 50 to about 65 weight percent, based on the total weight of the solution. The alkali metal halide can be, for example, lithium halide, preferably lithium bromide, although the absorption solution can include other alkali metal halides, such as lithium chloride, lithium iodide, as well as mixtures of these. Further, the absorption solution can include lithium nitrate. Still further, the absorption solution can include other halides, such as zinc halides, which are particularly useful in high temperature applications (generally about 450°F and higher). The zinc halides can be present in amounts up to about 45 weight percent (for example, a solution comprising 45 wt.% zinc halide and 20 wt.% lithium bromide).

As the skilled artisan will appreciate, the environment or conditions, such as temperature and/or pressure, of different refrigeration machines can vary. Typically, the temperature of the machine ranges from about 150°F to about 500°F, although the temperature can be outside this range as well. The solutions of the invention are particularly advantageous in higher temperature applications.

The present invention will be further illustrated by the following non-limiting examples.

EXAMPLE 1

Tests were performed in a pressure vessel autoclave with temperature control (+/- 2°F). A concentrated (65 wt.%) lithium bromide solution was prepared using anhydrous lithium bromide and deionized

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water. Sample solutions were prepared by adding an additive to the lithium bromide solution. Each additive was added individually to 800 ml lithium bromide solution. Pre-weighed metal coupons were 5 placed in and above the lithium bromide solution contained in a 2 liter cylinder made of Inconel-600 which fits into the autoclave. The cylinder containing the absorption solution and metal coupons was evacuated (to about 29.6 in of Hg) using a vacuum pump prior to 10 starting the test. The solution was heated to a specified temperature and held at that temperature for a period of 72 hours.

Upon completion, the test coupons were removed and cleaned. The corrosion rate was calculated 15 from the weight loss. The coupons were also tested by surface analysis. The results are set forth in Table 1 below. Corrosion rate (mpy) refers to mils per year.

TABLE I
Corrosion Rates for Carbon Steel at 72 Hours in 65 wt.% LiBr

	Additive	Concentration (ppm)	Corrosion Rate (mpy) at		
			350°F	400°F	450°F
20	None	--	41	220	326
	Lithium molybdate	160	64	38	43
	8-Hydroxyquinoline	500	4.5	6.5	37
		1000	5.0	5.8	13
		1500	0.7	1.7	4.8
		3000	0.5	0.7	2.0
	Irgacor-252	500	21	21	--
25	Durad 125	500	--	13	15
	Durad 220X	500	--	6.4	12
	Ortholum-162	500	--	9.6	16

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EXAMPLE 2

Tests were performed as described above in Example 1, except that corrosion rates were evaluated for carbon steel coupons in 65 wt.% LiBr solution at temperatures from 300-450°F for a period of 168 hours. The results are set forth in Table 2 below.

TABLE 2
Corrosion Rates for Carbon Steel at 168 Hours in 65 wt.% LiBr

	Additive	Corrosion Rate (mpy) at			
		300°F	350°F	400°F	450°F
10	None	25	60	118	132
	Lithium molybdate				
	199 ppm, pH = 11-12	9	10	12	14
	8-hydroxyquinoline				
15	1000 ppm, pH = 11-12	0.4	1.7	2.9	4.7
	1000 ppm, pH = 8-10	2.9	3.2	3.2	10

Hydrogen generation was also calculated and is set forth below for carbon steel in 65 wt.% LiBr at 168 hours for various additives. These results are set forth in Table 3 below.

20 **TABLE 3**
Hydrogen Generation from Carbon Steel in 65 wt.% LiBr at 168 Hours

	Additive	Hydrogen Generation (mg/in ²) at			
		300°F	350°F	400°F	450°F
25	None	0.5	27	36	37
	Lithium molybdate				
	199 ppm, pH = 11-12	2.7	2	6.9	7.9
	8-hydroxyquinoline				
	1000 ppm, pH = 12	0.1	1.6	5.4	13
	1000 ppm, pH = 9	0.5	7.6	7.8	21

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Surface analysis of the coupons exposed to the 8-hydroxyquinoline (1000 ppm)/LiBr solution (65 wt. %) at 400°F for one week was also conducted using X-ray diffraction and scanning electron microscopy/energy dispersive spectrometry (SEM/EDS) techniques. The X-ray diffraction measurements were carried out on a wide angle diffractometer using CuK α radiation with a diffracted beam monochromator. Figure 1 presents the X-ray diffraction pattern of a specific area of the metal coupon where a protective coating was formed after an attack from the brine solution. The peaks seen in the spectra identify formation of magnetite as the major phase on the metal surface.

The SEM/EDS analysis of the specific coated site on the treated coupon surface is presented in Figure 2. This spectra indicates that the protective film also contains carbon and oxygen atoms which belong to the 8-hydroxyquinoline molecule.

These results demonstrate that in the presence of 8-hydroxyquinoline, a protective coating containing carbon originating from the additive and magnetite is formed at the sites where the corrosion action of the brine solution occurs. The formation of the protective coating resists further degradation of metal due to corrosion.

EXAMPLE 3

L.E. Tsygankova, et al., Zaschita Metallov, vol. 27, No. 2. pps. 344-347 (1991), proposed a method for coating carbon steel (St3) with 8-hydroxyquinoline. The article reports holding carbon steel coupons in 0.5% or 0.2% solutions of 8-hydroxyquinoline ("8-HQ") in water at 100°C for 50 hours. The 50 hour period was stated to be sufficient for greater than 90% film buildup.

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The corrosion rates after keeping the coated steel in 40-60% LiBr solution at 150°C for 100 hours were reported to be as follows:

8-HQ in water, wt. %	Corrosion Rate at 150°C (100 hours) g/(m ² ,h)
0.2	0.33
0.5	0.47

To demonstrate the improved benefits of the present invention, the inventors followed procedure set forth in this article. Carbon steel coupons were 10 treated in a 0.5% solution of 8-HQ in water at 100°C for 50 hours. The coupon showed a non-uniform coating with blobs of material, several layers thick, isolated on the coupon surface. The coating was found to be gooey.

15 In a second test, three carbon steel coupons were treated in a 0.2% solution of 8-HQ in water at 100°C for 67 hours. A uniform film was formed on the coupon.

Scanning electron micrograph analysis was 20 conducted of the carbon steel coupons coated from solutions of 0.2 and 0.5 wt.% 8-HQ in water after the treatment with 8-HQ. The scan of the coupon treated with 0.5 wt.% 8-HQ solution showed a large deposition of the organic compound, as indicated by a large carbon 25 peak. Peaks for iron were very small, suggesting a morphology of a scale type, several layers thick, deposition of the organic compound.

The scan of the coupon with a relatively thinner layer of coating after treatment with 0.2 wt.% 30 8-HQ solution also showed a significant level of organic deposition, as indicated by a large carbon peak. Some areas also showed iron. Another scan of a cleaner area of this same coupon with relatively thinner coating showed some carbon deposition and a

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peak for the surface iron. This suggested a relatively thinner coating on the coupon.

Corrosion tests were also conducted of the resultant coupons. The coated coupons were prepared as 5 described in this example above in a 0.2 wt.% solution of 8-HQ in water at 100°C for 67 hours. The coupons were then air dried and tested in 65 wt.% LiBr solution at 400°F under vacuum of 29.6 inches of mercury.

Results are summarized below.

10 Corrosion rate (mpy) = 201, 223

Hydrogen evolved (mg/in²) = 59

The corrosion rate seen in this case was significantly high. It is also supported by the high level of hydrogen generation. After the test, the 15 coupon showed significant corrosion on the surface.

The L.E. Tsygankova, et al. article determined all corrosion rates reported therein at 150°C (302°F). The reported corrosion rate at 302°F for a 0.2% solution treated coupon was about 14.6 mpy. 20 Corrosion rate usually found at 300°F is relatively low.

In the present studies, the corrosion rate for carbon steel at 300°F in uninhibited 65 wt.% LiBr solution (0.06 N alkalinity) was found to be 25 mpy. 25 However, as demonstrated in Examples 1 and 2 above, the use of 8-HQ in situ in LiBr solutions resulted in remarkably low corrosion rates of less than 5 mpy under harsher conditions (450°F).

The above demonstrates that coating carbon 30 steel with 8-HQ as reported by the L.E. Tsygankova, et al. article does not provide adequate corrosion protection for LiBr. In comparison, the process of the invention carried out in LiBr in situ using 8-HQ have shown excellent corrosion protection at temperatures up 35 to 450°F.

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The foregoing examples are illustrative of the present invention and are not to be construed as limiting thereof.

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THAT WHICH IS CLAIMED IS:

1. A process for inhibiting the corrosion of a refrigeration machine resulting from the presence of alkali metal halide absorbent solutions, comprising
5 circulating in a refrigeration machine an absorption solution and at least one organic compound in an amount sufficient to provide a corrosion inhibiting effect.

2. The process of Claim 1, wherein said at least one organic compound has corrosion inhibiting
10 properties in absorption refrigeration systems.

3. The process of Claim 1, further comprises prior to said circulating step the step of adding said at least one organic compound directly into said refrigeration system.

15 4. The process of Claim 1, further comprises prior to said circulating step the step of adding said at least one organic compound to said absorption solution and thereafter adding said absorption solution to said refrigeration system.

20 5. The process of Claim 4, further comprises the step of heating said absorption solution prior to the step of adding said at least one organic compound to said absorption solution.

25 6. The process of Claim 1, wherein said absorption fluid comprises at least one organic compound capable of inhibiting corrosion.

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7. The process of Claim 1, wherein said at least one organic compound is selected from the group consisting of nitrogen containing heterobicyclic organic compounds, thio-carboxylic acids containing heterocyclic substituents, organophosphates and mixtures thereof.

8. The process of Claim 7, wherein said at least one nitrogen containing heterobicyclic nitrogen containing organic compound is selected from the group consisting of quinoline, tetrahydroquinoline, isoquinoline, tetrahydroisoquinoline, quinazoline, quinoxaline, 2,3-benzopyrrole, carboxylic, aldehyde or hydroxy derivatives thereof, and mixtures thereof.

9. The process of Claim 8, wherein said at least one nitrogen containing heterocyclic organic compound comprises 8-hydroxyquinoline.

10. The process of Claim 7, wherein said at least one thio-carboxylic acid containing heterocyclic substituents comprises at least one compound selected from the group consisting of benzothiazolyl-thio-carboxylic acids, benzoxazolyl-thio-carboxylic acids, benzylimidazolyl-thio-carboxylic acids, and mixtures thereof.

11. The process of Claim 10, wherein said at least one thio-carboxylic acid containing heterocyclic substituents comprises (2-benzothiazolylthio)-butanedioic acid.

12. The process of Claim 7, wherein said at least one organophosphate is selected from the group consisting of substituted and unsubstituted triphenyl phosphates, alkylated phenyl phosphate esters, alkyl acid orthophosphates, and mixtures thereof.

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13. The process of Claim 11, wherein said substituted and unsubstituted triphenyl phosphates comprises trixylenyl phosphate, tricresyl phosphate, and mixtures thereof.

5 14. The process of Claim 1, wherein during said circulating step, said at least one organic compound forms a protective layer of magnetite on a surface within said machine.

10 15. The process of Claim 1, wherein during said circulating step said solution is exposed to temperatures ranging from about 150°F to about 550°F.

15 16. The process of Claim 1, wherein said at least one organic compound is present in said refrigeration system in an amount of about 100 to about 5000 ppm.

17. An absorption solution for refrigeration systems, comprising at least one alkali metal halide and at least one organic compound in an amount sufficient to provide a corrosion inhibiting effect.

20 18. The solution of Claim 17, wherein said at least one organic compound has corrosion inhibiting properties in absorption refrigeration systems.

25 19. The solution of Claim 17, wherein said at least one organic compound is selected from the group consisting of nitrogen containing heterobicyclic organic compounds, thio-carboxylic acid containing heterocyclic substituents, organophosphates and mixtures thereof.

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20. The solution of Claim 19, wherein said at least one nitrogen containing heterobicyclic nitrogen containing organic compound is selected from the group consisting of quinoline, tetrahydroquinoline, 5 isoquinoline, tetrahydroisoquinoline, quinazoline, quinoxaline, 2,3-benzopyrrole, carboxylic, aldehyde or hydroxy derivatives thereof, and mixtures thereof.

21. The solution of Claim 20, wherein said at least one nitrogen containing heterocyclic organic 10 compound comprises 8-hydroxyquinoline.

22. The solution of Claim 19, wherein said at least one thio-carboxylic acid containing heterocyclic substituents comprises at least one compound selected from the group consisting of 15 benzothiazolyl-thio-carboxylic acids, benzoaxazolyl-thio-carboxylic acids, benzylimidazolyl-thio-carboxylic acids, and mixtures thereof.

23. The solution of Claim 22, wherein said at least one thio-carboxylic acid containing 20 heterocyclic substituents comprises (2-benzothiazolylthio)-butanedioic acid.

24. The solution of Claim 19, wherein said at least one organophosphate is selected from the group 25 consisting of substituted and unsubstituted triphenyl phosphates, alkylated phenyl phosphate esters, alkyl acid orthophosphates, and mixtures thereof.

25. The solution of Claim 24, wherein said substituted and unsubstituted triphenyl phosphates 30 comprises trixylenyl phosphate, tricresyl phosphate, and mixtures thereof.

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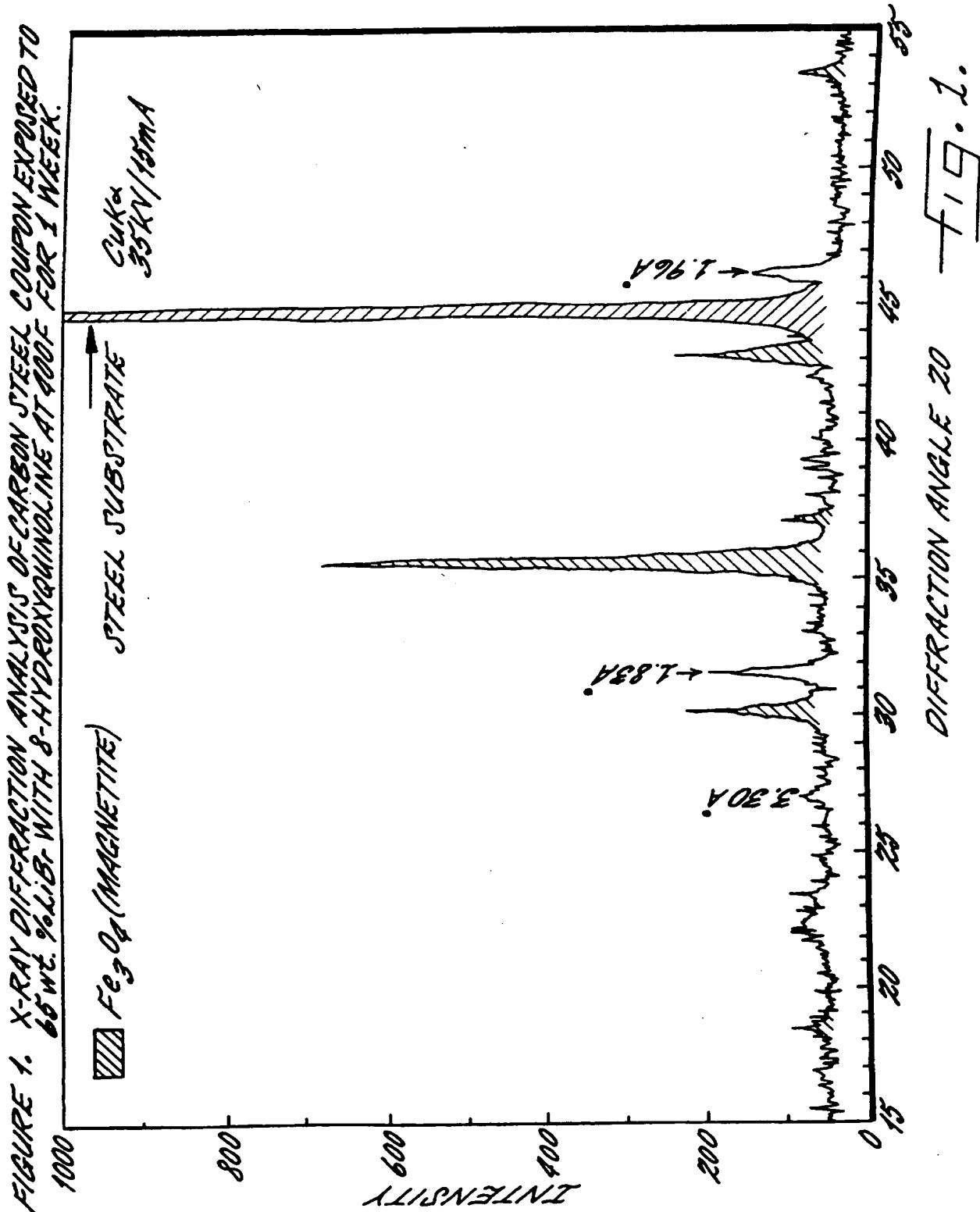
26. The solution of Claim 17, wherein said at least one organic compound is present in an amount of about 100 to about 5000 ppm.

27. The solution of Claim 17, wherein said
5 at least one alkali metal halide is selected from the group consisting of lithium bromide, lithium chloride, lithium iodide and mixtures thereof.

28. The solution of Claim 17, further comprising zinc halide.

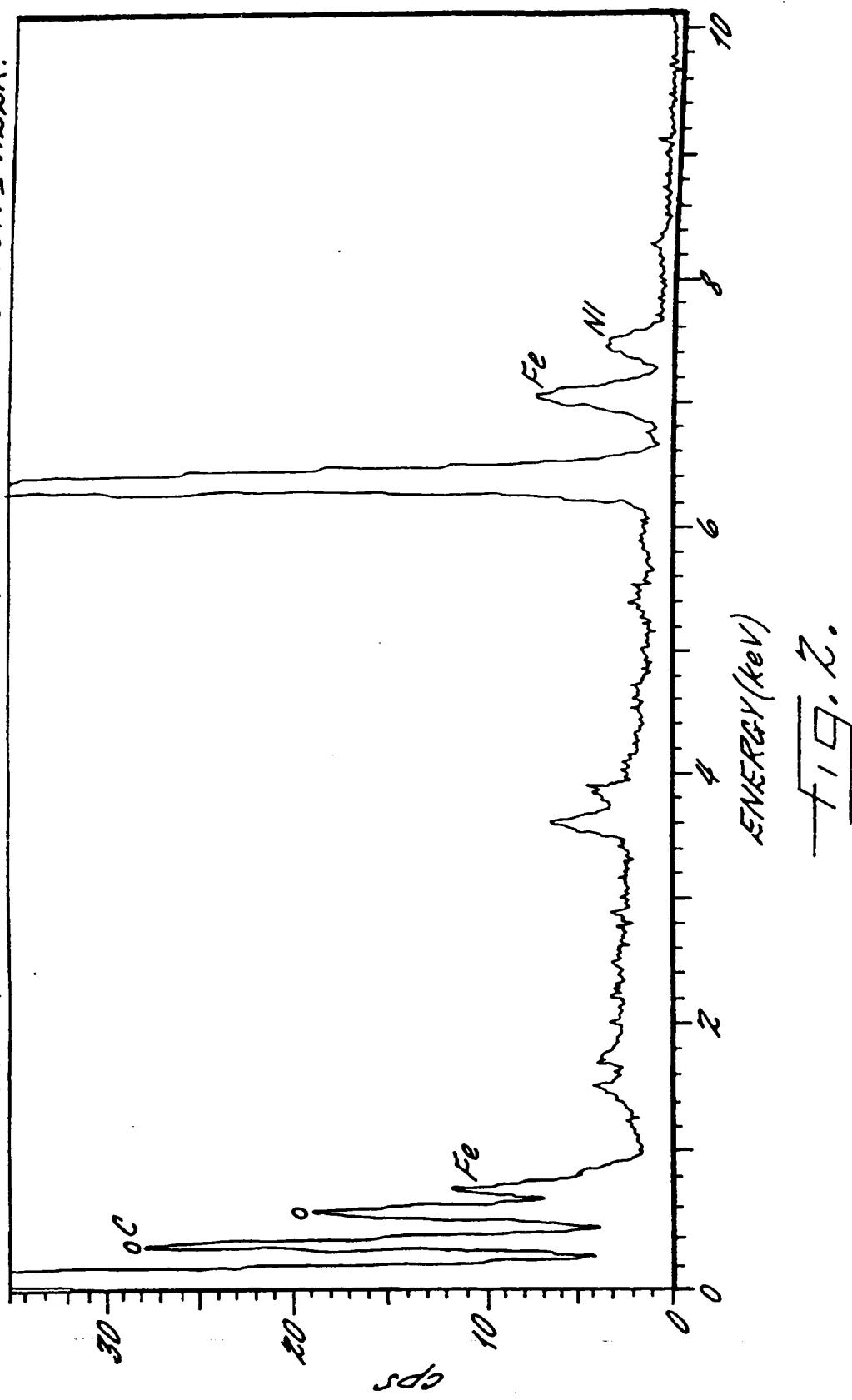
10 29. The solution of Claim 17, further comprising lithium nitrate.

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FIGURE 2. SEM/EDS ANALYSIS OF CARBON STEEL COUPON EXPOSED TO
65wt. % LiBr WITH 8-HYDROXYQUINOLINE AT 400F FOR 1 WEEK.



SUBSTITUTE SHEET (RULE 26)

INTERNATIONAL SEARCH REPORT

In International Application No.

PCT/US 97/13717

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6	C09K5/04	C23F11/14	C23F11/16
		C23F11/167	

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 C09K C23F C13F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	TSYGANKOVA L E ET AL: "PROTECTION OF STEEL BY HYDROXYQUINOLINE FILMS FROM CORROSION IN LITHIUM BROMIDE SOLUTIONS" PROTECTION OF METALS, vol. 27, no. 2, 1 March 1991, pages 288-291, XP000261669 cited in the application see the whole document	1-4,6-9, 15
Y	---	10-13, 20,21
X	US 4 311 024 A (ITOH MASAHIKO ET AL) 19 January 1982	1,2,4,6, 7,15-19, 26,27,29
Y	see the whole document	8-13, 20-25
	---	-/-

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Date of the actual completion of the international search

11 November 1997

Date of mailing of the international search report

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INTERNATIONAL SEARCH REPORT

Int'l Application No

PCT/US 97/13717

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>DATABASE WPI Week 8930 Derwent Publications Ltd., London, GB; AN 89-217080 XP002046273 & JP 01 155 157 A (SANYO ELECTRIC CO) , 19 June 1989 see abstract</p> <p>---</p>	1,2,4,6, 15-18,27
Y	<p>US 5 152 929 A (BENTLEY ROBERT L ET AL) 6 October 1992 cited in the application see the whole document</p> <p>---</p>	10,11, 22,23
Y	<p>EP 0 573 082 A (FMC CORP) 8 December 1993 cited in the application see the whole document</p> <p>---</p>	12,13, 24,25
Y	<p>SASTRI V S ET AL: "Surface analysis in corrosion inhibition mechanisms" WERKSTOFFE UND KORROSION, FEB. 1987, WEST GERMANY, vol. 38, no. 2, ISSN 0043-2822, pages 77-82, XP002046271 see page 80</p> <p>---</p>	8,9,20, 21
Y	<p>GRANESE S L ET AL: "The inhibition action of heterocyclic nitrogen organic compounds on Fe and steel in HCl media" CORROSION SCIENCE, SEPT. 1992, UK, vol. 33, no. 9, ISSN 0010-938X, pages 1439-1453, XP002046386 see table 1</p> <p>---</p>	8,20
Y	<p>US 4 563 253 A (LEIDHEISER JR HENRY ET AL) 7 January 1986 see table 1</p> <p>---</p>	8,9,20, 21
P,Y	<p>CHEMICAL ABSTRACTS, vol. 126, no. 3, 20 January 1997 Columbus, Ohio, US; abstract no. 34170, GORSKI, WIESLAW ET AL: "Brake fluid" XP002046390 see abstract</p> <p>& PL 166 099 A (WOJSKOWY OSRODEK BADAWCZO-ROZWOJOWY SLUZBY MATERIALOW PEDNYCH I SMAROW)</p> <p>---</p>	12,13, 24,25
5	<p>US 4 077 929 A (ROBINSON LARRY GRANT) 7 March 1978 see the whole document</p> <p>---</p>	12,13, 24,25
		-/-

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 97/13717

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	TANNO K ET AL: "The corrosion of carbon steel in lithium bromide solution at moderate temperatures" CORROSION SCIENCE, SEPT. 1993, UK, vol. 34, no. 9, ISSN 0010-938X, pages 1441-1451, XP002046272 see the whole document -----	14



INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 97/13717

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 4311024 A	19-01-82	JP 55085864 A	28-06-80
US 5152929 A	06-10-92	AU 563813 B AU 2794484 A CA 1225404 A DK 235384 A,B, EP 0129506 A JP 1737518 C JP 4025954 B JP 59212478 A US 5347008 A	23-07-87 15-11-84 11-08-87 15-11-84 27-12-84 26-02-93 06-05-92 01-12-84 13-09-94
EP 0573082 A	08-12-93	US 5206404 A IL 105410 A JP 2026306 C JP 6009661 A JP 7039429 B KR 9607100 B	27-04-93 18-02-97 26-02-96 18-01-94 01-05-95 27-05-96
US 4563253 A	07-01-86	US 4681814 A	21-07-87
US 4077929 A	07-03-78	DE 2609261 A	16-09-76